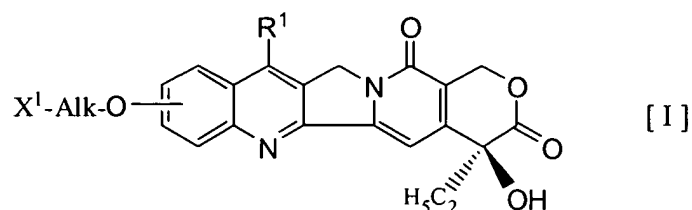


AMENDMENTS TO THE CLAIMS

Claims 1-19 (Cancelled)

20. (New) A liquid preparation consisting essentially of:

(a) 1 w/v% to 20 w/v% of a camptothecin derivative which is prepared by binding a compound of the formula [I]:



wherein R¹ is a substituted or unsubstituted lower alkyl group, X¹ is a group of the formula: -NHR² (wherein R² is a hydrogen atom or a lower alkyl group) or a hydroxyl group and Alk is a straight or branched chain alkylene group optionally interrupted by an oxygen atom, and a polysaccharide having carboxyl groups via an amino acid or a peptide, or a pharmaceutically acceptable salt thereof,

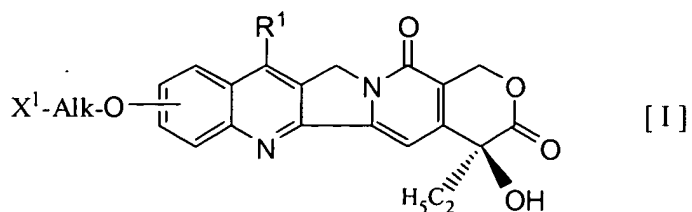
(b) a buffer and

(c) water,

which is adjusted to pH 5 - 8 with said buffer.

21. (New) A liquid preparation consisting essentially of:

(a) 1 w/v% to 20 w/v% of a camptothecin derivative which is prepared by binding a compound of the formula [I]:



wherein R^1 is a substituted or unsubstituted lower alkyl group, X^1 is a group of the formula: $-NHR^2$ (wherein R^2 is a hydrogen atom or a lower alkyl group) or a hydroxyl group and Alk is a straight or branched chain alkylene group optionally interrupted by an oxygen atom, and a polysaccharide having carboxyl groups via an amino acid or a peptide, or a pharmaceutically acceptable salt thereof,

(b) one or more stabilizers selected from an alkali metal carbonate and an alkali metal hydrogencarbonate,

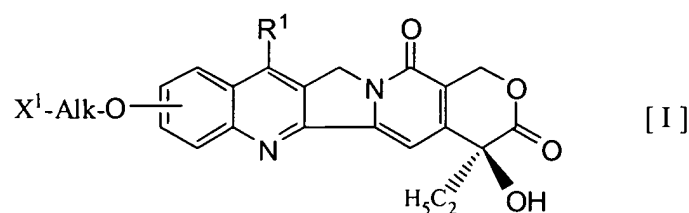
(c) a buffer comprising one or more compounds selected from the group consisting of citric acid, an alkali metal citrate, acetic acid, an alkali metal acetate and an alkali metal dihydrogenphosphate and

(d) water,

which is adjusted to pH 5 - 8 with said buffer.

22. (New) A liquid preparation consisting essentially of:

(a) 1 w/v% to 20 w/v% of a camptothecin derivative which is prepared by binding a compound of the formula [I]:



wherein R^1 is a substituted or unsubstituted lower alkyl group, X^1 is a group of the formula: $-NHR^2$ (wherein R^2 is a hydrogen atom or a lower alkyl group) or a hydroxyl group and Alk is a straight or branched chain alkylene group optionally interrupted by an oxygen atom, and a polysaccharide having carboxyl groups via an amino acid or a peptide, or a pharmaceutically acceptable salt thereof,

(b) one or more salts selected from the group consisting of an alkali metal chloride, an alkali earth metal chloride and an alkali metal sulphate,

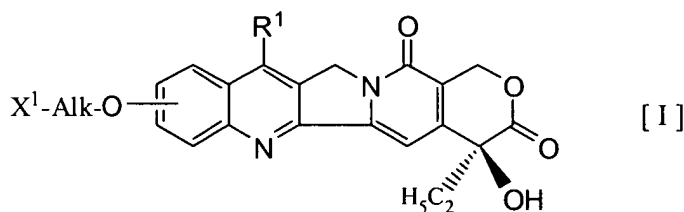
(c) a buffer comprising one or more compounds selected from the group consisting of citric acid, an alkali metal citrate, acetic acid, an alkali metal acetate and an alkali metal dihydrogenphosphate and

(d) water,

which is adjusted to pH 5 - 8 with said buffer.

23. (New) A liquid preparation consisting essentially of:

(a) 1 w/v% to 20 w/v% of a camptothecin derivative which is prepared by binding a compound of the formula [I]:



wherein R¹ is a substituted or unsubstituted lower alkyl group, X¹ is a group of the formula: -NHR² (wherein R² is a hydrogen atom or a lower alkyl group) or a hydroxyl group and Alk is a straight or branched chain alkylene group optionally interrupted by an oxygen atom, and a polysaccharide having carboxyl groups via an amino acid or a peptide, or a pharmaceutically acceptable salt thereof,

(b) one or more salts selected from the group consisting of an alkali metal chloride, an alkali earth metal chloride and an alkali metal sulphate,

(c) one or more stabilizers selected from an alkali metal carbonate and an alkali metal hydrogencarbonate,

(d) a buffer comprising one or more compounds selected from the group consisting of citric acid, an alkali metal citrate, acetic acid, an alkali metal acetate and an alkali metal dihydrogenphosphate and

(e) water,

which is adjusted to pH 5 - 8 with said buffer.

CCCCNC(=O)Oc1ccc2c(c1)c3nc4c2c(c3)nc5c4c(=O)oc(C[C@H](O)C)c5

30. (New) The liquid preparation according to claim 23, wherein the salt is sodium chloride.

31. (New) A lyophilized drug composition which is prepared by lyophilizing the liquid preparation claimed in claim 20.

32. (New) A lyophilized drug composition which is prepared by lyophilizing the liquid preparation claimed in claim 21.

33. (New) A lyophilized drug composition which is prepared by lyophilizing the liquid preparation claimed in claim 22.

34. (New) A lyophilized drug composition which is prepared by lyophilizing the liquid preparation claimed in claim 23.